Antiretroviral therapy in a community clinic — early lessons from a pilot project

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Objectives. To report on operational and clinical problems encountered during the first 6 months of a community-based antiretroviral therapy (ART) programme.

Methods. ART was implemented in a primary care setting utilising an easily replicable service-delivery model based on a medical officer and nurse. Therapeutic counsellors, themselves HIV-infected, provided counselling and adherence support. Drug and monitoring costs were charitably funded and provincial health authorities supplied the medical infrastructure. The HIV Research Unit, University of Cape Town, supplied training and additional clinical support. Local HIV primary care clinics provided patient referrals. Standardised ART regimens were used with strict entry criteria (AIDS or CD4 count < 200 cells/μl).

Results. Demand for the service was high. Referred patients had advanced disease (AIDS 57%, median CD4 count 96/μl) and high pre-treatment mortality (83/100 person-years). Mycobacterial disease was a major contributor to this mortality (40%). Scheduled clinic visit hours were six times higher during recruitment than maintenance. Attributable costs were: drugs 61%, staff 27%, viral load and CD4 cell counts 10% and safety monitoring 2%. Viral load after 16 weeks of therapy was < 400 copies/ml in the first 16 patients.

Conclusions. ART can be successfully implemented within a primary care setting. Drug purchases and staff salaries drive programme costing. The service model is capable of managing 250 - 300 patients on chronic ART, but staffing needs to be increased during recruitment. Attention must be given to the diagnosis of tuberculosis during screening and early ART. Incorporating therapeutic counsellors into the programme increased community involvement and utilised a valuable and previously untapped resource.
In South Africa, AIDS is now the major cause of death among young adults. While ART is available to thousands of South Africans in the private medical sector, there is little or no ART access in the public sector, despite demands for such therapy from the HIV-infected community. It has been argued that "where HIV is the leading cause of death, a basic minimum package that does not include antiretrovirals is not worthy of the name." 

In this paper we report on the initial medical, ethical and logistic challenges of initiating a community-based ART pilot project in a district where there is an existing network of HIV primary health care (PHC) clinics. The project was called Usapho Lwethu (My Family), since treatment was offered to individuals and their partners and children wherever appropriate. An easily replicable service delivery model was chosen based on a team consisting of a medical officer, nurse and therapeutic counsellors. In order to minimise operational complexity, standardised ART regimens were used, which complied with international and national treatment guidelines. To facilitate expansion of this or similar service delivery models for ART elsewhere in South Africa, there is urgent need to report and share knowledge of the operational challenges that need to be overcome.

**Methods**

The ART clinic was situated in the Guguletu Community Health Centre in the Nyanga district of Cape Town, which has a population of 325,436 (projected figure from 1996 census data). There are 10 primary care HIV clinics within the district, which served as the patient referral base. Each clinic had trained staff able to provide voluntary testing and counselling services together with prophylaxis and outpatient management of common HIV opportunistic infections. Patients requiring inpatient care were referred to a local 200-bed secondary hospital.

Funding for antiretroviral medication, viral load measurements and CD4 cell counts, sufficient for 150 patients, was provided by UK-based charities. The local Community Health Service Organisation and AIDS Directorate of the Western Cape provided clinic space and medical support services, together with dedicated clinic staff, consisting of a medical officer and nurse. Training, clinical support, protocol development and programme evaluation were provided by the HIV Research Unit of the University of Cape Town. In August 2002, medical staff of the referring PHC clinics were invited to workshops, where eligibility criteria and patient referral mechanisms were discussed. Referring doctors were requested to select suitable drug-naive candidates from their regular clinic attendees and to fax contact details and a medical summary to the programme nurse. The medical criteria for ART eligibility were either a prior AIDS diagnosis or a CD4 cell count less than 200 cells/μl.

Branded medications registered with the Medicines Control Council of South Africa were used in standard doses. ART consisted of two schedules, an initial non-nucleoside reverse transcriptase (NNRTI)-based regimen of stavudine (d4T), lamivudine (3TC) and efavirenz (nevirapine was substituted for efavirenz in women of child-bearing potential), with a second protease inhibitor (PI)-based regimen of zidovudine (AZT), didanosine (ddI) and Kaletra. Medication was sourced from a single pharmaceutical supplier with a rapid delivery turnaround time, to minimise on-site drug stocks. Drugs were packaged for each named patient and supplied in 4 x 30 day packs and stored in a secure locked section of the main clinic pharmacy.

During September 2002, the first patients were recruited into the programme, with a schedule of visits as follows: -4 weeks, -2 weeks, treatment initiation, +4 weeks, +8 weeks, +16 weeks and then at 16-week intervals. The early schedule of visits was truncated for women in the latter stages of pregnancy, to maximise therapeutic benefit to mother and baby. At the screening visit (week -4), potential candidates were allocated a therapeutic counsellor, whose responsibility was to provide ongoing counselling support, to reinforce the need for high levels of adherence, to maintain communication between clients and the clinic staff and to visit clients in their homes. The therapeutic counsellors were HIV-infected individuals, many on ART themselves, who had been trained in drug adherence and ART toxicity recognition by a non-governmental organisation, Scapila. The therapeutic counsellor/patient ratio was 1:20. At this initial visit, a treatment readiness assessment questionnaire was completed and 4 weeks of co-trimoxazole were dispensed, with pill counts performed after 14 and 28 days to assess adherence. All patients had a checklist completed, which screened for symptoms and signs of tuberculosis (TB) infection. At the week -2 visit, blood was taken for quantitative viral load and CD4 cell count. These tests were repeated at 16-week intervals thereafter. Drug safety monitoring of liver function test (LFT) and full blood count (FBC) were also performed at weeks -2, +8 and 16-weekly thereafter, with an additional LFT performed at week +2 in those receiving nevirapine to screen for potential hepatotoxicity. The final decision to commence or defer initiation of ART was made at a combined meeting of medical and therapeutic counsellors during the third week (week -1), when information on clinical status, blood tests, treatment-readiness and adherence data were available. Following this meeting, drugs were ordered for those commencing ART. The cohort pre-ART period was defined as the cumulative number of days between the screening visit and the date of either commencement of ART or permanent deferral. The ART exposure period was defined as the cumulative number of days from ART commencement to the censoring date of

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28 February 2003. Resource utilisation data including scheduled and non-scheduled visits and staff allocated time were prospectively collected in a clinic register. Clinical, laboratory and adherence data were kept in an electronic database.

Results

The number of eligible patients referred and subsequent status within the programme are outlined in Fig. 1. The mean age of those screened was 34 years (range 7 - 55 years); 78% were female. The population had advanced HIV infection, with 95% having symptomatic disease (WHO clinical stages 3 and 4) and 57% having AIDS (WHO clinical stage 4). The median CD4 count was 96 cells/μl (range 3 - 452 cells/μl), and the median viral load was 4.91 logs (range 2.611 - 500.000).

Eight patients were permanently deferred; 4 because of failure or inability to attend clinic visits, 1 because of advanced debilitating HIV disease and multidrug-resistant pulmonary tuberculosis, 2 because of asymptomatic HIV and CD4 cell counts above entry criteria at screening, and 1 because he denied being HIV-infected. One patient was withdrawn after therapy commenced because of failure to attend clinic visits. Viral load data at 16 weeks were available for the first 16 patients, all of which were < 400 copies/ml. The first child born to a mother on the programme was HIV PCR-negative at 3 months.

Five deaths occurred between screening and the commencement of ART or treatment deferral. The total pre-ART period for the cohort was 2 186 days, which translated to a pre-treatment crude mortality rate of 83/100 patient years. Two of the five patients had active Mycobacterium tuberculosis infection. Four deaths occurred on ART, none of which was directly related to ART. The period of ART exposure up to the censoring date totalled 4 271 days.

The proportional costs of drugs, staffing, quantitative viral load with CD4 cell counts and laboratory safety monitoring are shown in Fig. 2. Programme costs were dominated by drug procurement and personnel costs. Of the personnel costs, the medical officer contributed 39.3%, the nurse 16%, Sizophila counsellors 15.7% and specialist medical support and training 29%.

Financial costing for the first 6 months of HAART project

Drug 27%
Staff 61%
Monitoring 3%
Laboratory safety 9%

Fig. 2. The first 6 months' financial costs of the programme introducing ART to HIV primary care service. The attributable proportion of costs for the following components are shown: drug acquisition; staffing — medical, nursing and counsellors; monitoring of CD4 and viral load; and laboratory safety monitoring.

Patient referrals reached 150 after 8 weeks of the programme. Initiation of the 150 patients onto ART was scheduled over 9 months, with a mean of 15 patients starting therapy per month. In the first 6 months there were 421 scheduled and 20 unscheduled visits. Duration of scheduled visits varied between 15 minutes for a week-8 visit and 45 minutes for a treatment-initiation visit. The number of scheduled clinic hours per week was not evenly distributed and the projected weekly scheduled visit hours required for the programme are shown in Fig. 3.

Discussion

This ART pilot project is a unique collaboration between funding organisations from a developed country, local and
allocate increased logistical support to ART clinics during the initial 'set-up' phase.

Patients on the programme had a functional status sufficient to be able to attend an outpatient clinic; however, the pre-treatment crude mortality rate was 83/100 patient years. The on-treatment mortality showed significant decline; the mortality rate has not been presented at this stage because the drug exposure period is subject to a major confounder of 'right censoring' and these data will be reported later. TB was a significant contributor to mortality and the exclusion of active TB presents a particular challenge to those initiating ART in high TB-prevalence populations such as South Africa.\textsuperscript{13,15}

Present treatment guidelines\textsuperscript{16} give an upper threshold of 200 CD4 cells/μl for initiating therapy, but there is no lower limit. This introduces the dilemma of whether patients are ever too sick to join a programme. While the sickest patients have much to gain, they utilise programme resources disproportionately and a high death rate can impact negatively on community perceptions of the programme and affect staff morale adversely. The non-medical therapeutic counsellors required specific counselling support around this issue. In an analysis of 12 574 individuals on ART, the CD4 count and clinical stage at the time of initiation of treatment were the dominant prognostic factors.\textsuperscript{14} In a cohort with very advanced HIV disease and low CD4 counts such as ours, between 11% and 15% would be predicted to develop further AIDS-related complications during the first year of therapy, with the majority of these events occurring within the first 6 months.\textsuperscript{16} Despite this high predicted complication rate, European and North American experience indicates that only 1.5 - 3.1% of advanced patients with 50 - 100 CD4 cells/μl die in the first year of ART.\textsuperscript{16} These data support initiation of therapy at low CD4 counts, which can still result in a reasonable prognosis if the initial complications around the time of starting therapy can be managed. Staff of the clinic and referral hospital therefore need a high level of expertise in the recognition and management of medical complications occurring during early ART, particularly if very ill patients are to be entered into a programme.

Drug procurement accounted for 61% of the attributable cost of adding ART to the existing primary care HIV management programme. Although retail costs of ART have declined in South Africa in recent years, they remain a major obstacle to wider implementation in the public sector. In the next few months the programme is expected to access UNAIDS preferential pricing which will reduce the cost of the initial regimen to approximately 50% of current retail pricing. Availability of generic formulations could reduce the cost of our initial schedule by an additional 50%, to approximately R350 per month.\textsuperscript{14,17} Medical specialist support contributed 29% of initial staffing costs; however, this component will also decrease as training requirements decline and an increasing

Fig. 3. Number of hours of scheduled visits per week required for 150 patients initiating HAART over a 9-month period. There is an initial staffing requirement, which peaks at approximately six times the long-term staffing needs.
but also on the allocation of sufficient resources for the sites will be dependent not only on funding for antiretrovirals, counsellors gave valuable input to the programme from inclusion and exclusion of very sick patients was very challenging and stressful for the clinic team. Therapeutic counsellors gave valuable input to the programme from persons in the local community living with HIV and AIDS. The incorporation of HIV-infected individuals as 'treatment buddies' and to give counselling support meant that a previously untapped resource of expertise and talent was utilised.

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References